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(54) Title: COUPLING OF BLENDS OF α -OLEFIN/VINYL AROMATIC MONOMER OR HINDERED ALIPHATIC VINYL MONOMER INTERPOLYMERS WITH POLYOLEFINS (57) Abstract The invention includes a process of preparing a coupled polymer blend by heating an admixture containing (1) a polymer blend of (A) from 1 to 99 weight percent of one or more α -olefin/hindered vinyl monomer substantially random interpolymers, each having been made from monomer components comprising: (1) from 0.5 to 65 mole percent of either (a) at least one vinyl aromatic monomer, or (b) at least one hindered aliphatic vinyl monomer, or (c) a combination of at least one vinyl aromatic monomer and at least one hindered aliphatic vinyl monomer; and (2) from 35 to 99.5 mole percent of at least one aliphatic α -olefin having from 2 to 20 carbon atoms; and (B) from 99 to 1 weight percent of one or more homopolymers or copolymers made from monomer components comprising aliphatic α -olefins having from 2 to 20 carbon atoms, or aliphatic α -olefins having from 2 to 20 carbon atoms and containing polar groups; and (2) a coupling amount of at least one poly(sulfonyl azide) to at least the decomposition temperature of the poly(sulfonyl azide).		

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COUPLING OF BLENDS OF α -OLEFIN/VINYL AROMATIC MONOMER OR HINDERED
ALIPHATIC VINYL MONOMER INTERPOLYMERS WITH POLYOLEFINS

The present invention pertains to blends of interpolymers made from monomer components comprising at least one α -olefin and at least
5 one aromatic vinyl or vinylidene monomer or at least one hindered aliphatic vinyl or vinylidene monomer or at least one cycloaliphatic vinyl or vinylidene monomer with olefinic polymers, or a combination thereof especially such blends which are coupled.

Interpolymers of the class which include α -olefin/hindered
10 vinyl monomer substantially random interpolymers and include materials such as α -olefin/vinyl aromatic monomer interpolymers are known in the art and offer a range of material structures and properties which makes them useful for varied applications, such as compatibilizers for blends of polyethylene and polystyrene as
15 described in US 5,460,818.

As described by D'Anniello et al. (Journal of Applied Polymer Science, Volume 58, pages 1701-1706 (1995)), such interpolymers can show good elastic properties and energy dissipation characteristics. Alternatively, certain of the interpolymers are useful in adhesive
20 systems, as illustrated for instance in United States patent number 5,244,996, issued to Mitsui Petrochemical Industries Ltd.

It would be desirable to provide improvements in processability or performance without the addition of additives commonly used to improve properties of the interpolymers or enhance properties that
25 can be achieved by using additives and would be more desirable to achieve combinations of properties not attainable using blends of polymers alone. Alternatively, characteristics of a blend produced to obtain certain desirable characteristics would advantageously be further improved by practice of the present invention.
30 Characteristics which would desirably improve would include at least one of low temperature toughness, mechanical strength, heat resistance and melt processability, preferably heat resistance.

Coupling of polymers results in rheology modification of the polymer. As used herein, the term "rheology modification" means
35 change in melt viscosity of a polymer as determined by dynamic mechanical spectroscopy. Preferably the melt strength increases while maintaining the high shear viscosity (that is viscosity measured at a shear of 100 rad/sec by DMS) so that a polymer exhibits more resistance to stretching during elongation of molten polymer at
40 low shear conditions (that is viscosity measured at a shear of 0.1 rad/sec by DMS) and does not sacrifice the output at high shear conditions.

Use of poly(sulfonyl azide)s to react with polymers is known for instance in The teachings of US 3,058,944; 3,336,268; and 3,530,108 include the reaction of certain poly(sulfonyl azide) compounds with isotactic polypropylene or other polyolefins by
5 nitrene insertion into C-H bonds. The product reported in US 3,058,944 is crosslinked. The product reported in US 3,530,108 is foamed and cured with cycloalkane-di(sulfonyl azide) of a given formula. In US 3,336,268 the resulting reaction products are referred to as "bridged polymers" because polymer chains are
10 "bridged" with sulfonamide bridges. The disclosed process includes a mixing step such as milling or mixing of the sulfonylazide and polymer in solution or dispersion then a heating step where the temperature is sufficient to decompose the sulfonylazide (100°C to 225° depending on the azide decomposition temperature). The starting
15 polypropylene polymer for the claimed process has a molecular weight of at least 275,000. Blends taught in US 3,336,268 have up to 25 percent ethylene propylene elastomer.

Similarly, the teachings of Canadian patent 797,917 (family member of NL 6,503,188) include rheology modification using from
20 0.001 to 0.075 weight percent polysulfonyl azide to modify homopolymer polyethylene and its blend with polyisobutylene.

It would also be desirable to use a polymer of enhanced melt strength in a foaming process preferably to achieve at least one of smaller cell diameter, homogeneous cell diameter distribution, lower
25 foam density, higher tensile and compressive strength, higher tensile or compressive toughness. Preferably, the polymer would have at most little high shear viscosity increase over a corresponding polymer of the same chemical composition, but not modified to obtain the enhanced melt strength, for example the starting material polymer, so
30 that it would have similar processing characteristics as the unmodified polymer.

It has been found that blends of interpolymers made from monomer components comprising at least one α -olefin and at least one aromatic vinyl monomer or at least one hindered aliphatic vinyl
35 monomer or at least one cycloaliphatic vinyl monomer with olefinic polymers, or a combination thereof said blends having been heated with at least one poly(sulfonyl azide) at least to the decomposition temperature of the poly(sulfonyl azide), advantageously have improvements in at least one of low temperature toughness, mechanical
40 strength, heat resistance or processability, preferably heat resistance.

This improvement is believed to result from the poly(sulfonyl azide) reacting with more than one polymer chain to connect them, referred to herein as "coupling" and the poly(sulfonyl azide), as a "coupling agent." Coupling results in and is measured by rheology
5 modification.

Preferably, blends of the invention having their rheology modified by treatment with poly(sulfonyl azide) coupling agents advantageously have increased melt strength useful in foaming.

The invention includes a process of preparing a coupled polymer
10 blend comprising heating an admixture containing (1) a polymer blend containing:

(A) from 1 to 99 weight percent based on entire composition of one or more α -olefin/hindered vinyl monomer substantially random interpolymers, each having been made from monomer components comprising:

- 15 (1) from 0.5 to 65 mole percent of either
(a) at least one vinyl aromatic monomer, or
(b) at least one hindered aliphatic vinyl monomer, or
(c) a combination of at least one vinyl aromatic monomer and at least one hindered aliphatic monomer; and

- 20 (2) from 35 to 99.5 mole percent of at least one aliphatic α -olefin having from 2 to 20 carbon atoms; and

(B) from 99 to 1 weight percent of one or more homopolymers or copolymers made from monomer components comprising aliphatic α -olefins having from 2 to 20 carbon atoms, or aliphatic α -olefins having from 2
25 to 20 carbon atoms and containing polar groups; and (2) a coupling amount of at least one poly(sulfonyl azide) to at least the decomposition temperature of the poly(sulfonyl azide) for a period sufficient for decomposition of at least 80 weight percent of the poly(sulfonyl azide) and sufficient to result in a coupled polymer blend
30 having less than 2 weight percent gel. Preferably component (A) is a substantially random interpolymer made from styrene and ethylene, or styrene, ethylene and at least one other α -olefin containing from 3 to 8 carbon atoms; and component (B) is a homopolymer made from ethylene or propylene, or a copolymer made from ethylene, or propylene, or a
35 combination thereof and at least other α -olefin containing from 4 to 8 carbon atoms; or a terpolymer of ethylene, propylene and at least one of 4-methyl pentene, butene-1, hexene-1 or octene-1. The poly(sulfonyl azide) and blend preferably react at a temperature which is at least greater than the first temperature, at least the decomposition
40 temperature of the poly(sulfonyl azide) and greater than 150°C. More

preferably the poly(sulfonyl azide) and blend are mixed at a first temperature which is at least the melting point of the lowest melting component of the blend and after mixing react at a second temperature which is at least greater than the first temperature, is at least the decomposition temperature of the poly(sulfonyl azide) and is greater than 185°C.

Further, the invention includes any composition formed by a process of the invention, especially a composition comprising a reaction product obtainable by heating an admixture containing (1) a polymer blend containing:

(A) from 1 to 99 weight percent of one or more α -olefin/hindered vinyl monomer substantially random interpolymers, each having been made from monomer components comprising:

(1) from 0.5 to 65 mole percent of either

- (a) at least one vinyl aromatic monomer, or
- (b) at least one hindered aliphatic vinyl monomer, or
- (c) a combination of at least one vinyl aromatic monomer and at least one hindered aliphatic vinyl monomer; and

(2) from 35 to 99.5 mole percent of at least one aliphatic α -olefin having from 2 to 20 carbon atoms; and

(B) from 99 to 1 weight percent of one or more homopolymers or copolymers made from monomer components comprising aliphatic α -olefins having from 2 to 20 carbon atoms, or aliphatic α -olefins having from 2 to 20 carbon atoms and containing polar groups; and (2) a coupling amount at least one poly(sulfonyl azide) to at least the decomposition temperature of the poly(sulfonyl azide) for a period sufficient for decomposition of at least 80 weight percent of the poly(sulfonyl azide).

The invention also includes any an article which comprises a composition of the invention. Preferably the article is formed from a melt of the composition. More preferably it is formed by a melt process, most preferably the article is calendared, cast and blown sheet, film, compression and injection molded part, fiber, modifier for bitumen or asphalt compositions, or a component in a hot melt or pressure sensitive adhesive system or foam.

The term "interpolymer" is used herein to indicate a polymer wherein at least two different monomers are polymerized to make the interpolymer.

The term "copolymer" as employed herein means a polymer wherein at least two different monomers are polymerized to form the copolymer.

The term "mer(s)" means the polymerized unit of the polymer derived from the indicated monomer(s).

The term "monomer residue" or "polymer units derived from" means that portion of the polymerizable monomer molecule which
5 resides in the polymer chain as a result of being polymerized with another polymerizable molecule to make the polymer chain.

The term "substantially random" in the substantially random interpolymer resulting from polymerizing one or more α -olefin monomers and one or more vinyl or vinylidene aromatic monomers or
10 hindered aliphatic or cycloaliphatic vinyl or vinylidene monomers, and optionally, with other polymerizable ethylenically unsaturated monomer(s) as used herein means that the distribution of the monomers of said interpolymer can be described by the Bernoulli statistical model or by a first or second order Markovian statistical model, as
15 described by J. C. Randall in POLYMER SEQUENCE DETERMINATION, Carbon-13 NMR Method, Academic Press New York, 1977, pp. 71-78. Preferably, the substantially random interpolymer resulting from polymerizing one or more α -olefin monomers and one or more vinyl aromatic monomer, and optionally, with other polymerizable ethylenically unsaturated
20 monomer(s) does not contain more than 15 percent of the total amount of vinyl aromatic monomer in blocks of vinyl aromatic monomer of more than 3 units. More preferably, the interpolymer was not characterized by a high degree of either isotacticity or syndiotacticity. This means that in the carbon-¹³ NMR spectrum of the
25 substantially random interpolymer the peak areas corresponding to the main chain methylene and methine carbons representing either meso diad sequences or racemic diad sequences should not exceed 75 percent of the total peak area of the main chain methylene and methine carbons.

Any numerical values recited herein include all values from the
30 lower value to the upper value in increments of one unit provided that there is a separation of at least 2 units between any lower value and any higher value. As an example, if it is stated that the amount of a component or a value of a process variable such as, for
35 example, temperature, pressure, or time is, for example, from 1 to 90, preferably from 20 to 80, more preferably from 30 to 70, it is intended that values such as 15 to 85, 22 to 68, 43 to 51, 30 to 32 etc. are expressly enumerated in this specification. For values which are less than one, one unit is considered to be 0.0001, 0.001,
40 0.01 or 0.1 as appropriate. These are only examples of what is specifically intended and all possible combinations of numerical values between the lowest value and the highest value enumerated are

to be considered to be expressly stated in this application in a similar manner.

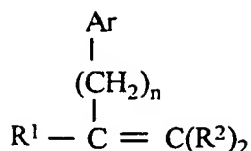
Polymers used as starting materials in the practice of the invention are blends of polymers. Each polymer in the blend is a polyolefin. At least one of the polyolefins is interpolymers made from monomer components comprising at least one α -olefin and at least one aromatic vinyl monomer or at least one hindered aliphatic vinyl monomer or at least one cycloaliphatic vinyl monomer or a combination thereof (A) and at least one other polymer is a different polyolefin (B). There is preferably a phase separated polymer composition containing distinct hard and soft segments, where the hard segments reinforce the soft phase, but it is not crosslinked into a network, that is the composition is still thermoplastic. In blends there are advantageously improvements of at least one physical property such as impact strength, stiffness, heat resistance, low temperature toughness, mechanical strength, scratch and mar resistance, or processability of these blends as compared to blends of the same components not treated by the process of the invention.

The interpolymers suitable as component (A) for the blends comprising the present invention include substantially random interpolymers prepared by polymerizing one or more α -olefin monomers with one or more vinyl aromatic monomers, or one or more hindered aliphatic or cycloaliphatic vinyl monomers, or a combination thereof, and optionally with other polymerizable ethylenically unsaturated monomer(s).

Suitable α -olefin monomers include for example, α -olefin monomers containing from 2 to 20, preferably from 2 to 12, more preferably from 2 to 8 carbon atoms. Preferred such monomers include ethylene, propylene, butene-1, 4-methyl-1-pentene, hexene-1 and octene-1. Most preferred are ethylene or a combination of ethylene with C_{3-8} α -olefins. These α -olefins do not contain an aromatic moiety.

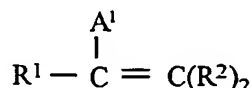
Other optional polymerizable ethylenically unsaturated monomer(s) include strained ring olefins such as norbornene and C_{1-10} alkyl or C_{6-10} aryl substituted norbornenes, with an exemplary interpolymer being ethylene/styrene/norbornene.

Suitable vinyl aromatic monomers which can be employed to prepare the interpolymers employed in the blends include, for example, those represented by the following formula:



wherein R¹ is selected from the group of radicals consisting of hydrogen and alkyl radicals containing from 1 to 4 carbon atoms, preferably hydrogen or methyl; each R² is independently selected from the group of radicals consisting of hydrogen and alkyl radicals containing from 1 to 4 carbon atoms, preferably hydrogen or methyl; Ar is a phenyl group or a phenyl group substituted with from 1 to 5 substituents selected from the group consisting of halo, C₁₋₄-alkyl, and C₁₋₄-haloalkyl; and n has a value from zero to 6, preferably from zero to 2, most preferably zero. Exemplary vinyl aromatic monomers include styrene, vinyl toluene, α-methylstyrene, t-butyl styrene, chlorostyrene, including all isomers of these compounds. Particularly suitable such monomers include styrene and lower alkyl- or halogen-substituted derivatives thereof. Preferred monomers include styrene, α-methyl styrene, the lower alkyl- (C₁ - C₄) or phenyl-ring substituted derivatives of styrene, such as for example, ortho-, meta-, and para-methylstyrene, the ring halogenated styrenes, para-vinyl toluene or mixtures thereof. A more preferred aromatic vinyl monomer is styrene.

By the term "hindered aliphatic or cycloaliphatic vinyl or vinylidene compounds", it is meant addition polymerizable vinyl or vinylidene monomers corresponding to the formula:



wherein A¹ is a sterically bulky, aliphatic or cycloaliphatic substituent of up to 20 carbons, R¹ is selected from the group of radicals consisting of hydrogen and alkyl radicals containing from 1 to 4 carbon atoms, preferably hydrogen or methyl; each R² is independently selected from the group of radicals consisting of hydrogen and alkyl radicals containing from 1 to 4 carbon atoms, preferably hydrogen or methyl; or alternatively R¹ and A¹ together form a ring system. By the term "sterically bulky" is meant that the monomer bearing this substituent is normally incapable of addition polymerization by standard Ziegler-Natta polymerization catalysts at a rate comparable with ethylene polymerizations. Preferred aliphatic or cycloaliphatic vinyl or vinylidene monomers are those in which one of the carbon atoms bearing ethylenic unsaturation is tertiary or quaternary substituted. Examples of such substituents include cyclic

aliphatic groups such as cyclohexyl, cyclohexenyl, cyclooctenyl, or ring alkyl or aryl substituted derivatives thereof, tert-butyl, and norbornyl. Most preferred hindered aliphatic or cycloaliphatic vinyl or vinylidene compounds are the various isomeric vinyl- ring substituted derivatives of cyclohexene and substituted cyclohexenes, and 5-ethylidene-2-norbornene. Especially suitable are 1-, 3-, and 4-vinylcyclohexene.

The interpolymers of one or more α -olefins and one or more vinyl aromatic monomers one or more hindered aliphatic or cycloaliphatic vinyl monomers employed in the present invention are substantially random polymers. These interpolymers usually contain from 0.5 to 65, preferably from 1 to 55, more preferably from 2 to 50 mole percent of at least one vinyl aromatic monomer hindered aliphatic or cycloaliphatic vinyl monomer and from 35 to 99.5, preferably from 45 to 99, more preferably from 50 to 98 mole percent of at least one aliphatic α -olefin having from 2 to 20 carbon atoms.

Other optional polymerizable ethylenically unsaturated monomer(s) include strained ring olefins such as norbornene and C_{1-10} alkyl or C_{6-10} aryl substituted norbornenes, with an exemplary interpolpolymer being ethylene/styrene/norbornene.

The number average molecular weight (Mn) of the polymers and interpolymers is usually greater than 5,000, preferably from 20,000 to 1,000,000, more preferably from 50,000 to 500,000.

Polymerizations and unreacted monomer removal at temperatures above the autopolymerization temperature of the respective monomers may result in formation of some amounts of homopolymer polymerization products resulting from free radical polymerization. For example, while preparing the substantially random interpolpolymer, an amount of atactic vinyl aromatic homopolymer may be formed due to homopolymerization of the vinyl aromatic monomer at elevated temperatures. The presence of vinyl aromatic homopolymer is in general not detrimental for the purposes of the present invention and can be tolerated. The vinyl aromatic homopolymer may be separated from the interpolpolymer, if desired, by extraction techniques such as selective precipitation from solution with a non solvent for either the interpolpolymer or the vinyl aromatic homopolymer. For the purpose of the present invention it is preferred that no more than 20 weight percent, preferably less than 15 weight percent based on the total weight of the interpolymers of vinyl aromatic homopolymer is present.

The substantially random interpolymers are optionally modified by typical grafting, hydrogenation, functionalizing, or other reactions well known to those skilled in the art. The polymers may

be readily sulfonated or chlorinated to provide functionalized derivatives according to established techniques.

The substantially random interpolymers are prepared by polymerizing a mixture of polymerizable monomers in the presence of metallocene or constrained geometry catalysts for instance as described in to EP-A-0,416,815 by James C. Stevens et al. and U.S. Patent No. 5,703,187 by Francis J. Timmers. Preferred operating conditions for such polymerization reactions are pressures from atmospheric up to 3,000 atmospheres and temperatures from -30°C to 200°C.

Examples of suitable catalysts and methods for preparing the substantially random interpolymers are also disclosed in U.S. Application No. 07/702,475, filed May 20, 1991 corresponding to EP-A-514,828; as well as U.S. Patents: 5,055,438; 5,057,475; 5,096,867; 5,064,802; 5,132,380; 5,189,192; 5,321,106; 5,347,024; 5,350,723; 5,374,696; 5,399,635; and 5,721,185.

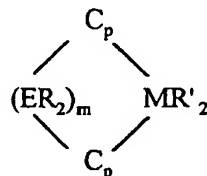
The substantially random α -olefin/vinyl aromatic interpolymers can also be prepared by the methods described by John G. Bradfute et al. (W. R. Grace & Co.) in WO 95/32095; by R. B. Pannell (Exxon Chemical Patents, Inc.) in WO 94/00500; and in Plastics Technology, p. 25 (September 1992).

Also suitable are the substantially random interpolymers which comprise at least one α -olefin/vinyl aromatic/vinyl aromatic/ α -olefin tetrad disclosed in U. S. Application No. 08/708,809 filed September 4, 1996 by Francis J. Timmers et al. These interpolymers contain additional signals with intensities greater than three times the peak to peak noise. These signals appear in the chemical shift range 43.70-44.25 ppm and 38.0-38.5 ppm. Specifically, major peaks are observed at 44.1, 43.9 and 38.2 ppm. A proton test NMR experiment indicates that the signals in the chemical shift region 43.70-44.25 ppm are methine carbons and the signals in the region 38.0-38.5 ppm are methylene carbons.

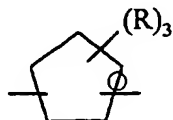
In order to determine the carbon-¹³ NMR chemical shifts of the interpolymers described, the following procedures and conditions are employed. A five to ten weight percent polymer solution is prepared in a mixture consisting of 50 volume percent 1,1,2,2-tetrachloroethane-d₂ and 50 volume percent 0.10 molar chromium tris(acetylacetonate) in 1,2,4-trichlorobenzene. NMR spectra are acquired at 130°C using an inverse gated decoupling sequence, a 90° pulse width and a pulse delay of five seconds or more. The spectra are referenced to the isolated methylene signal of the polymer assigned at 30.000 ppm.

It is believed that these new signals are due to sequences involving two head-to-tail vinyl aromatic monomer preceded and followed by at least one α -olefin insertion, for example an ethylene/styrene/styrene/ethylene tetrad wherein the styrene monomer
 5 insertions of said tetrads occur exclusively in a 1,2 (head to tail) manner. It is understood by one skilled in the art that for such tetrads involving a vinyl aromatic monomer other than styrene and an α -olefin other than ethylene that the ethylene/vinyl aromatic
 10 monomer/vinyl aromatic monomer/ethylene tetrad will give rise to similar carbon-¹³ NMR peaks but with slightly different chemical shifts.

These interpolymers are prepared by conducting the polymerization at temperatures of from -30°C to 250°C in the presence of such catalysts as those represented by the formula



15 wherein: each C_p is independently, each occurrence, a substituted cyclopentadienyl group π -bound to M; E is C or Si; M is a group IV metal, preferably Zr or Hf, most preferably Zr; each R is independently, each occurrence, H, hydrocarbyl, silahydrocarbyl, or
 20 hydrocarbylsilyl, containing up to 30 preferably from 1 to 20 more preferably from 1 to 10 carbon or silicon atoms; each R' is independently, each occurrence, H, halo, hydrocarbyl, hydrocarbyloxy, silahydrocarbyl, hydrocarbylsilyl containing up to 30 preferably from 1 to 20 more preferably from 1 to 10 carbon or silicon atoms or two
 25 R' groups together can be a C_{1-10} hydrocarbyl substituted 1,3-butadiene; m is 1 or 2; and optionally, but preferably in the presence of an activating cocatalyst. Particularly, suitable substituted cyclopentadienyl groups include those illustrated by the formula:



30 wherein each R is independently, each occurrence, H, hydrocarbyl, silahydrocarbyl, or hydrocarbylsilyl, containing up to 30 preferably from 1 to 20 more preferably from 1 to 10 carbon or silicon atoms or two R groups together form a divalent derivative of such group.

Preferably, R independently each occurrence is (including where appropriate all isomers) hydrogen, methyl, ethyl, propyl, butyl, pentyl, hexyl, benzyl, phenyl or silyl or (where appropriate) two such R groups are linked together forming a fused ring system such as
5 indenyl, fluorenyl, tetrahydroindenyl, tetrahydrofluorenyl, or octahydrofluorenyl.

Particularly preferred catalysts include, for example, racemic-(dimethylsilanediyl(2-methyl-4-phenylindenyl))zirconium dichloride, racemic-(dimethylsilanediyl(2-methyl-4-phenylindenyl))zirconium 1,4-
10 diphenyl-1,3-butadiene, racemic-(dimethylsilanediyl(2-methyl-4-phenylindenyl))zirconium di-C₁-, alkyl, racemic-(dimethylsilanediyl(2-methyl-4-phenylindenyl))zirconium di-C₁-, alkoxide, or any combination thereof.

Further preparative methods for the interpolymer component (A)
15 of the present invention have been described in the literature. Longo and Grassi (Makromol. Chem., Volume 191, pages 2387 to 2396 [1990]) and D'Anniello et al. (Journal of Applied Polymer Science, Volume 58, pages 1701-1706 [1995]) reported the use of a catalytic system based on methylalumoxane (MAO) and cyclopentadienyltitanium trichloride
20 (CpTiCl₃) to prepare an ethylene-styrene copolymer. Xu and Lin (Polymer Preprints, Am.Chem.Soc., Div.Polym.Chem.) Volume 35, pages 686,687 [1994]) have reported copolymerization using a MgCl₂/TiCl₄/NdCl₃/Al(iBu)₃ catalyst to give random copolymers of styrene and propylene. Lu et al (Journal of Applied Polymer Science,
25 Volume 53, pages 1453 to 1460 [1994]) have described the copolymerization of ethylene and styrene using a TiCl₄/NdCl₃/ MgCl₂ /Al(Et)₃ catalyst. Sernetz and Mulhaupt, (Macromol. Chem. Phys., v. 197, pp 1071-1083, 1997) have described the influence of polymerization conditions on the copolymerization of styrene with
30 ethylene using Me₂Si(Me,Cp)(N-tert-butyl)TiCl₄/methylaluminoxane Ziegler-Natta catalysts. The manufacture of α -olefin/vinyl aromatic monomer interpolymers such as propylene/styrene and butene/styrene are described in United States patent number 5,244,996, issued to Mitsui Petrochemical Industries Ltd. .

35 Olefinic polymers suitable for use as component (B) in the blends according to the present invention are aliphatic α -olefin homopolymers or interpolymers, or interpolymers of one or more aliphatic α -olefins and one or more non-aromatic monomers interpolymerizable therewith such as C₂-C₂₀ α -olefins or those
40 aliphatic α -olefins having from 2 to 20 carbon atoms and containing polar groups. Suitable aliphatic α -olefin monomers which introduce

polar groups into the polymer include, for example, ethylenically unsaturated nitriles such as acrylonitrile, methacrylonitrile, ethacrylonitrile, etc.; ethylenically unsaturated anhydrides such as maleic anhydride; ethylenically unsaturated amides such as acrylamide, methacrylamide etc.; ethylenically unsaturated carboxylic acids (both mono- and difunctional) such as acrylic acid and methacrylic acid, etc.; esters (especially lower, for example C₁-C₄, alkyl esters) of ethylenically unsaturated carboxylic acids such as methyl methacrylate, ethyl acrylate, hydroxyethylacrylate, n-butyl acrylate or methacrylate, 2-ethyl-hexylacrylate etc.; ethylenically unsaturated dicarboxylic acid imides such as N-alkyl or N-aryl maleimides such as N-phenyl maleimide, etc. Preferably such monomers containing polar groups are acrylic acid, vinyl acetate, maleic anhydride and acrylonitrile. Halogen groups which can be included in the polymers from aliphatic α -olefin monomers include fluorine, chlorine and bromine; preferably such polymers are chlorinated polyethylenes (CPEs). Preferred olefinic polymers for use in the present invention are homopolymers or interpolymers of an aliphatic, including cycloaliphatic, α -olefin having from 2 to 18 carbon atoms. Suitable examples are homopolymers of ethylene or propylene, and interpolymers of two or more α -olefin monomers. Other preferred olefinic polymers are interpolymers of ethylene and one or more other α -olefins having from 3 to 8 carbon atoms. Preferred comonomers include 1-butene, 4-methyl-1-pentene, 1-hexene, and 1-octene. The olefinic polymer blend component (B) may also contain, in addition to the α -olefin, one or more non-aromatic monomers interpolymerizable therewith. Such additional interpolymerizable monomers include, for example, C₄-C₂₀ dienes, preferably, butadiene or 5 ethylidene-2-norbornene. The olefinic polymers can be further characterized by their degree of long or short chain branching and the distribution thereof.

One class of olefinic polymers is generally produced by a high pressure polymerization process using a free radical initiator resulting in the traditional long chain branched low density polyethylene (LDPE). LDPE employed in the present composition usually has a density of less than 0.94 g/cc (ASTM D 792) and a melt index of from 0.01 to 100, and preferably from 0.1 to 50 grams per 10 minutes (as determined by ASTM Test Method D 1238, condition 190°/2.16).

Another class is the linear olefin polymers which have an absence of long chain branching, as the traditional linear low

density polyethylene polymers (heterogeneous LLDPE) or linear high density polyethylene polymers (HDPE) made using Ziegler polymerization processes (for example, U.S. Patent No. 4,076,698 (Anderson et al.), sometimes called heterogeneous polymers.

5 HDPE consists mainly of long linear polyethylene chains. The HDPE employed in the present composition usually has a density of at least 0.94 grams per cubic centimeter (g/cc) as determined by ASTM D 792, and a melt index (ASTM-1238, condition 190°/2.16) in the range of from 0.01 to 100, and preferably from 0.1 to 50 grams per 10
10 minutes.

The heterogeneous LLDPE employed in the present composition generally has a density of from 0.85 to 0.94 g/cc (ASTM D 792), and a melt index (ASTM-1238, condition 190°/2.16) in the range of from 0.01 to 100, and preferably from 0.1 to 50 grams per 10 minutes.
15 Preferably the LLDPE is an interpolymer of ethylene and one or more other α -olefins having from 3 to 18 carbon atoms, more preferably from 3-8 carbon atoms. Preferred comonomers include 1-butene, 4-methyl-1-pentene, 1-hexene, and 1-octene.

A further class is that of the uniformly branched or
20 homogeneous polymers (for example, homogeneous polyethylene). The homogeneous polymers contain no long chain branches and have only branches derived from the monomers (if having more than two carbon atoms). Homogeneous polymers include those made as described in U.S. Patent 3,645,992 (Elston), and those made using so-called single site
25 catalysts in a batch reactor having relatively high olefin concentrations (as described in U.S. Patent Nos. 5,026,798 and 5,055,438 (Canich). The uniformly branched/homogeneous polymers are those polymers in which the comonomer is randomly distributed within a given interpolymer molecule and wherein the interpolymer molecules
30 have a similar ethylene/comonomer ratio within that interpolymer.

The homogeneous LLDPE employed in the present composition generally has a density of from 0.85 to 0.94 g/cc (ASTM D 792), and a melt index (ASTM-1238, condition 190°/2.16) in the range of from 0.01 to 100, and preferably from 0.1 to 50 grams per 10 minutes.
35 Preferably the LLDPE is an interpolymer of ethylene and one or more other α -olefins having from 3 to 18 carbon atoms, more preferably from 3-8 carbon atoms. Preferred comonomers include 1-butene, 4-methyl-1-pentene, 1-hexene, and 1-octene.

Further, there is the class of substantially linear olefin
40 polymers that may advantageously be used in component (B) of the blends of the present invention. These polymers have a

processability similar to LDPE, but the strength and toughness of LLDPE. Substantially linear olefin polymers are disclosed in U.S. Patent Nos. 5,272,236 and 5,278,272.

The density of the substantially linear olefin polymers as
5 measured in accordance with ASTM D-792 is generally from 0.85 g/cc to 0.97 g/cc, preferably from 0.85 g/cc to 0.955 g/cc, and especially from 0.85 g/cc to 0.92 g/cc.

The melt index, according to ASTM D-1238, Condition 190°C/2.16 kg (also known as I₂), of the substantially linear olefin polymer is
10 generally from 0.01 g/10 min. to 1000 g/10 min., preferably from 0.01 g/10 min. to 100 g/10 min., and especially from 0.01 g/10 min. to 10 g/10 min.

Also, included are the ultra low molecular weight ethylene polymers and ethylene/ α -olefin interpolymers described in the U. S. Patent Application No. 784,683 entitled Ultra-Low Molecular Weight
15 Polymers, filed January 22, 1997 M. J. Guest, et al.. These ethylene/ α -olefin interpolymers have I₂ melt indices greater than 1,000, or a number average molecular weight (Mn) less than 11,000.

The substantially linear olefin polymer can be a homopolymer of
20 C₂-C₂₀ olefins, such as ethylene, propylene, 4-methyl-1-pentene, etc., or it can be an interpolymer of ethylene with at least one C₃-C₂₀ α -olefin, or C₂-C₂₀ acetylenically unsaturated monomer, or C₄-C₁₈ diolefin, or a combination thereof. The polymer can also be an interpolymer of ethylene with at least one of the above C₃-C₂₀ α -
25 olefins, diolefins, or acetylenically unsaturated monomers, or a combination thereof with additional other unsaturated monomers.

Especially preferred olefin polymers suitable for use as component (B) comprise LDPE, HDPE, heterogeneous LLDPE, homogeneous linear polyethylene, substantially linear olefin polymer,
30 polypropylene (PP), especially isotactic polypropylene and rubber toughened polypropylenes, or ethylene-propylene interpolymers (EP), or chlorinated polyolefins (CPE), or ethylene-vinyl acetate copolymers, or ethylene-acrylic acid copolymers, or any combination thereof. More preferably, at least one of the polymers (B) is a
35 propylene polymer or a homo- or co-polymer of ethylene with at least one other addition polymerizable monomer, most preferably a propylene polymer, a high density polyethylene, an ethylene α olefin copolymer having a heat resistance (melting point) greater than 90°C, or ethylene vinyl acetate copolymer (EVA), particularly a propylene
40 polymer, that is a polymer having at least 50 mole percent propylene repeating units. Ethylene/ α -olefin copolymers are preferably at

least 40 weight percent ethylene. Polypropylene is useful to improve heat resistance and modulus of Component A. Heat resistance of Component A is also conveniently improved by blending with high density polyethylene, ethylene alpha-olefin copolymer with a

- 5 crystalline melting point (T_m) greater than 90°C or a mixture thereof. An ethylene alpha-olefin copolymer with a crystalline melting point (T_m) less than 80°C is useful to blend with Component A to improve the low temperature toughness thereof. Alternatively, Component B is optionally an ethylene vinyl acetate copolymer (EVA)
- 10 to improve polarity of Component A.

- Advantageously, the blends of the present invention comprise from 1 to 99, preferably from 5 to 95 and more preferably from 10 to 90 percent by weight of the interpolymers containing at least one aromatic vinyl monomer residue or hindered aliphatic or
- 15 cycloaliphatic vinyl monomer residue or any combination thereof (component (A)) and from 1 to 99, preferably from 5 to 95, more preferably from 10 to 90 percent by weight of the polymers which do not contain any aromatic vinyl monomer residue or hindered aliphatic or cycloaliphatic vinyl monomer residue (component (B)). For high
- 20 heat elastomers, the blend is preferably at least 80 and more preferably no more than 60 weight percent Component (A) and preferably at least 20, more preferably at most 60 weight percent Component (B). For thermoplastic applications like automotive parts, appliances, building and construction materials, and films, the blend
- 25 is preferably at least 10 and more preferably no more than 30 weight percent Component (A) and preferably at least 70, more preferably at most 90 weight percent Component (B). The percentages are based on the total amount of the polymers constituting the blends.

- The blends of the present invention are prepared by any
- 30 suitable means within the skill in the art such as, but not limited to, dry blending in a pelletized form in the desired proportions followed by melt blending in a screw extruder, Banbury mixer or the like. Dry blended pellets are conveniently directly melt processed into a final solid state article by for example injection molding.
- 35 Alternatively, the blends are optionally made by direct polymerization, without isolation of the blend components, using for example one or more catalysts in one reactor or two or more reactors in series or parallel.

- For the purposes of coupling and, optionally, foaming, the
- 40 polymer is reacted with a polyfunctional compound capable of insertion reactions into C-H bonds. Such polyfunctional compounds have at least two, preferably 2, functional groups capable of C-H

insertion reactions. Those skilled in the art are familiar with C-H insertion reactions and functional groups capable of such reactions. For instance, carbenes as generated from diazo compounds, as cited in Mathur, N.C.; Snow, M.S.; Young, K.M., and Pincock, J.A.;

- 5 Tetrahedron, (1985), 41(8), pages 1509-1516, and nitrenes as generated from azides, as cited in Abramovitch, R.A.,; Chellathurai, T.; Holcomb, W.D; McMaster, I.T.; and Vanderpool, D.P.; J. Org. Chem., (1977), 42(17), 2920-6, and Abramovitch, R.A., Knaus, G.N., J. Org. Chem., (1975), 40(7), 883-9.

- 10 Compounds having at least two functional groups capable of C-H insertion under reaction conditions are referred to herein as coupling agents.

- Polyfunctional compounds capable of insertions into C-H bonds include poly(sulfonyl azide)s. The poly(sulfonyl azide) is any
15 compound having at least two sulfonyl azide groups ($-SO_2N_2$) reactive with the polyolefin. Preferably the poly(sulfonyl azide)s have a structure X-R-X wherein each X is SO_2N_2 and R represents an unsubstituted or inertly substituted hydrocarbyl, hydrocarbyl ether or silicon-containing group, preferably having sufficient carbon,
20 oxygen or silicon, preferably carbon, atoms to separate the sulfonyl azide groups sufficiently to permit a facile reaction between the polyolefin and the sulfonyl azide, more preferably at least 1, more preferably at least 2, most preferably at least 3 carbon, oxygen or silicon, preferably carbon, atoms between functional groups. While
25 there is no critical limit to the length of R, each R advantageously has at least one carbon or silicon atom between X's and preferably has less than 50, more preferably less than 30, most preferably less than 20 carbon, oxygen or silicon atoms. Within these limits, larger is better for reasons including thermal and shock stability. When R
30 is straight-chain alkyl hydrocarbon, there are preferably less than 4 carbon atoms between the sulfonyl azide groups to reduce the propensity of the nitrene to bend back and react with itself. Silicon containing groups include silanes and siloxanes, preferably siloxanes. The term inertly substituted refers to substitution with
35 atoms or groups which do not undesirably interfere with the desired reaction(s) or desired properties of the resulting coupled polymers. Such groups include fluorine, aliphatic or aromatic ether, siloxane as well as sulfonyl azide groups when more than two polyolefin chains are to be joined. Suitable structures include R as aryl, alkyl, aryl
40 alkaryl, arylalkyl silane, siloxane or heterocyclic, groups and other groups which are inert and separate the sulfonyl azide groups as described. More preferably R includes at least one aryl group between the sulfonyl groups, most preferably at least two aryl groups

(such as when R is 4,4' diphenylether or 4,4'-biphenyl). When R is one aryl group, it is preferred that the group have more than one ring, as in the case of naphthylene bis(sulfonyl azides).

Poly(sulfonyl)azides include such compounds as 1, 5-pentane
5 bis(sulfonlazide), 1,8-octane bis(sulfonyl azide), 1,10-decane
bis(sulfonyl azide), 1,10-octadecane bis(sulfonyl azide), 1-octyl-
2,4,6-benzene tris(sulfonyl azide), 4,4'-diphenyl ether bis(sulfonyl
azide), 1,6-bis(4'-sulfonazidophenyl)hexane, 2,7-naphthalene
bis(sulfonyl azide), and mixed sulfonyl azides of chlorinated
10 aliphatic hydrocarbons containing an average of from 1 to 8 chlorine
atoms and from 2 to 5 sulfonyl azide groups per molecule, and
mixtures thereof. Preferred poly(sulfonyl azide)s include oxy-bis(4-
sulfonylazidobenzene), 2,7-naphthalene bis(sulfonyl azido), 4,4'-
bis(sulfonyl azido)biphenyl, 4,4'-diphenyl ether bis(sulfonyl azide)
15 and bis(4-sulfonyl azidophenyl)methane, and mixtures thereof.

Sulfonyl azides are conveniently prepared by the reaction of sodium azide with the corresponding sulfonyl chloride, although oxidation of sulfonyl hydrazines with various reagents (nitrous acid, dinitrogen tetroxide, nitrosonium tetrafluoroborate) has been used.

20 Polyfunctional compounds capable of insertions into C-H bonds also include carbene-forming compounds such as salts of alkyl and aryl hydrazones and diazo compounds, and nitrene-forming compounds such as alkyl and aryl azides ($R-N_3$), acyl azides ($R-C(O)N_3$), azidoformates ($R-O-C(O)-N_3$), sulfonyl azides ($R-SO_2-N_3$), phosphoryl
25 azides ($(RO)_2-(PO)-N_3$), phosphinic azides ($R_2-P(O)-N_3$) and silyl azides (R_3-Si-N_3). Some of the coupling agents of the invention are preferred because of their propensity to form a greater abundance of carbon-hydrogen insertion products. Such compounds as the salts of hydrazones, diazo compounds, azidoformates, sulfonyl azides,
30 phosphoryl azides, and silyl azides are preferred because they form stable singlet-state electron products (carbenes and nitrenes) which carry out efficient carbon-hydrogen insertion reactions, rather than substantially 1) rearranging via such mechanisms as the Curtius-type rearrangement, as is the case with acyl azides and phosphinic azides,
35 or 2) rapidly converting to the triplet-state electron configuration which preferentially undergoes hydrogen atom abstraction reactions, which is the case with alkyl and aryl azides. Also, selection from among the preferred coupling agents is conveniently possible because of the differences in the temperatures at which the different classes
40 of coupling agents are converted to the active carbene or nitrene products. For example, those skilled in the art will recognize that carbenes are formed from diazo compounds efficiently at temperatures less than 100°C, while salts of hydrazones, azidoformates and the

sulfonyl azide compounds react at a convenient rate at temperatures above 100°C, up to temperatures of 200°C. (By convenient rates it is meant that the compounds react at a rate that is fast enough to make commercial processing possible, while reacting slowly enough to allow
5 adequate mixing and compounding to result in a final product with the coupling agent adequately dispersed and located substantially in the desired position in the final product. Such location and dispersion are sometimes different from product to product depending on the desired properties of the final product.) Phosphoryl azides are
10 conveniently reacted at temperatures in excess of 180°C up to 300°C, while silyl azides react preferentially at temperatures of from 250°C to 400°C.

The amount of poly(sulfonyl azide) used to treat blends by the practice of the invention is an amount sufficient to result in
15 rheology modification of the blend, preferably in improved properties of the blend or to reduce the average particle size of the dispersed phase as observed by electron microscopy as compared with a blend of the same components formed with the same mixing and some other conditions but without reaction with the poly(sulfonyl azide)
20 referred to herein as a "coupling amount." Properties improved preferably include at least one of low temperature toughness as measured by ASTM D 256-84, mechanical strength as measured by ASTM D-412, heat resistance as measured by a thermomechanical analyzer (TMA) (for which procedural detail is included hereinafter),
25 or processability as measured by a dynamic mechanical spectrometer (DMS) (for which procedural detail is included hereinafter.) To avoid the detrimental effects of gels, the amount is preferably less than the amount sufficient to result in 4, preferably 2 weight percent gels in the coupled blend. This amount is preferably at least 0.01,
30 more preferably at least 0.05 and preferably less than 0.5 more preferably less than 0.4, most preferably less than 0.3 weight percent of poly(sulfonyl azide) based on total weight of polymers in the blend.

For coupling, the sulfonyl azide is admixed with the polymer
35 and heated to at least the decomposition temperature of the sulfonyl azide. By decomposition temperature of the azide it is meant that temperature at which the azide converts to the sulfonyl nitrene, eliminating nitrogen and heat in the process, as determined by differential scanning calorimetry (DSC). The poly(sulfonyl
40 azide) begins to react at a kinetically significant rate (convenient for use in the practice of the invention) at temperatures of 130°C and is almost completely reacted at 160°C in a DSC (scanning at

10°C/min). The temperature of the peak observed in a DSC scan of decomposition v. temperature, is referred to herein as the "peak decomposition temperature" and is the decomposition temperature referred to herein unless stated otherwise. ARC (scanning at 2°C/hr) shows onset of decomposition is 100°C. Extent of reaction is a function of time and temperature. At the low levels of azide used in the practice of the invention, the optimal properties are not reached until the azide is essentially fully reacted. Temperatures for use in the practice of the invention are also determined by the softening or melt temperatures of the polymer starting materials. For these reasons, the temperature is advantageously greater than 90°C, preferably greater than 120°C, more preferably greater than 150°C, most preferably greater than 170°C. Formation of free radicals is preferably avoided; therefore, temperatures in excess of 250°C are preferably avoided; more preferably the temperature is less than 200°C.

Preferred times at the desired decomposition temperatures are times that are sufficient to result in reaction of the coupling agent with the polymer(s) without undesirable thermal degradation of the polymer matrix. Preferred reaction times in terms of the half life of the coupling agent, that is the time required for half of the agent to be reacted at a preselected temperature, which half life is determinable by DSC is 5 half lives of the coupling agent. In the case of a bis(sulfonyl azide), for instance, the reaction time is preferably at least 4 minutes at 200 °C.

Admixing of the polymer and coupling agent is conveniently accomplished by any means within the skill in the art. Desired distribution is different in many cases, depending on what properties are to be modified. In a blend it is often desirable to have low solubility in one or more of the polymer matrices such that the azide is preferentially in the other phase, or predominantly in the interfacial region between the two phases

Treatment of blends with the poly(sulfonyl azide) according to practice of the invention results in blends of the invention which are referred to herein as chain coupled, reactively coupled or coupled blends. A blend is advantageously mixed with a poly(sulfonyl azide) above the softening temperature of at least one component of the blend, most preferably below the peak decomposition temperature of the poly(sulfonyl azide), and the resulting mixture is preferably raised to at least the peak decomposition temperature of the poly(sulfonyl azide). Practice of the invention advantageously

involves forming a substantially uniform admixture of polymers and poly(sulfonyl azide) before decomposition of the poly(sulfonyl azide), although in the case of blends where there are dispersed and continuous phases, it is sufficient that the poly(sulfonyl azide) be
5 dispersed at the interface of the phases rather than uniformly distributed in particularly the dispersed phase unless chain coupling of the dispersed phase itself is desired. Most preferably, the poly(sulfonyl azide) and resulting coupling is distributed primarily at the interface of the different polymers. Distribution primarily
10 at the interface is advantageously achieved by adding the polyazide after the two immiscible polymers have been mixed to the extent that a minimum dispersed polymer particle size has been achieved. This allows for the maximum amount of interfacial surface area to be available for reaction of the polyazide.

15 Where there are dispersed and continuous phases, it is most preferable, but not necessary, to add the poly(sulfonyl azide) after the blend of two or more polymers is well mixed, that is at a point when the particle size of the dispersed polymer has reached the smallest size practically attainable on the particular mixing device.
20 being used. At least one of the blend polymer components is preferably at least at its softening temperature. More preferably mixing occurs or is continued when the blend is at a temperature sufficient for the poly(sulfonyl azide) to react to form a reactive species believed to be a singlet nitrene capable of inserting into
25 carbon-hydrogen bonds, that is at its decomposition temperature. This allows for optimum reaction at the interface between the two polymers. While it is preferred that mixing of the blend and poly(sulfonyl azide) precede a temperature increase to the decomposition temperature, alternatively, mixing occurs at or above
30 the decomposition temperature of the poly(sulfonyl azide).

It is believed that the improved properties exhibited by blends of the invention results from the formation of polymers coupled between components of the blend. This coupled polymer would then act as a compatibilizer and lower the interfacial tension between the
35 blend components. The result is believed to be a finer dispersion of the dispersed phase or a coupling of dispersed particles to the continuous phase polymer leading to improved properties. For use in foaming the coupled polymer is optionally coupled separately from the foaming process and used in a foaming process within the skill in the
40 art. Such processes for coupling include at least one of (a) dry blending the coupling agent with the polymer, preferably to form a substantially uniform admixture and adding this mixture to melt processing equipment, for example a melt extruder to achieve the

coupling reaction, at a temperature at least the decomposition temperature of the coupling agent; (b) introducing, for example by injection, a coupling agent in liquid form, for example dissolved in a solvent therefor or in a slurry of coupling agent in a liquid, into
5 a device containing polymer, preferably softened, molten or melted polymer, but alternatively in particulate form, in solution or dispersion, more preferably in melt processing equipment; (c) forming a first admixture of a first amount of a first polymer and a coupling agent, advantageously at a temperature less than the decomposition
10 temperature of the coupling agent, preferably by melt blending, and then forming a second admixture of the first admixture with a second amount of a second polymer (for example a concentrate of a coupling agent admixed with at least one polymer and optionally other additives, is conveniently admixed into a second polymer or
15 combination thereof optionally with other additives, to modify the second polymer(s)); (d) feeding at least one coupling agent, preferably in solid form, more preferably finely comminuted, for example powder, directly into softened or molten polymer, for example in melt processing equipment, for example in an extruder; or
20 combinations thereof. Among processes (a) through (d), processes (b) and (c) are preferred, with (c) most preferred. For example, process (c) is conveniently used to make a concentrate with a first polymer composition having a lower melting temperature, advantageously at a temperature below the decomposition temperature of the coupling
25 agent, and the concentrate is melt blended into a second polymer composition having a higher melting temperature to complete the coupling reaction. Concentrates are especially preferred when temperatures are sufficiently high to result in loss of coupling agent by evaporation or decomposition not leading to reaction with
30 the polymer, or other conditions would result that effect. Alternatively, some coupling occurs during the blending of the first polymer and coupling agent, but some of the coupling agent remains unreacted until the concentrate is blended into the second polymer composition. Each polymer or polymer composition includes at least
35 one homopolymer, copolymer, terpolymer, or interpolymer and optionally includes additives within the skill in the art. When the coupling agent is added in a dry form it is preferred to mix the agent and polymer in a softened or molten state below the decomposition temperature of the coupling agent then to heat the
40 resulting admixture to a temperature at least equal to the decomposition temperature of the coupling agent.

The term "melt processing" is used to mean any process in which the polymer is softened or melted, such as extrusion, pelletizing,

molding, thermoforming, film blowing, compounding in polymer melt form, or fiber spinning.

The polyolefin(s) and coupling agent are suitably combined in any manner which results in desired reaction thereof, preferably by
5 mixing the coupling agent with the polymer(s) under conditions which allow sufficient mixing before reaction to avoid uneven amounts of localized reaction then subjecting the resulting admixture to heat sufficient for reaction. Preferably, a substantially uniform admixture of coupling agent and polymer is formed before exposure to
10 conditions in which chain coupling takes place. A substantially uniform admixture is one in which the distribution of coupling agent in the polymer is sufficiently homogeneous to be evidenced by a polymer having a melt viscosity after treatment according to the practice of the invention either higher at low angular frequency (for
15 example 0.1 rad/sec) or lower at higher angular frequency (for example 100 rad/sec) than that of the same polymer which has not been treated with the coupling agent but has been subjected to the same shear and thermal history. Thus, preferably, in the practice of the invention, decomposition of the coupling agent occurs after mixing
20 sufficient to result in a substantially uniform admixture of coupling agent and polymer. This mixing is preferably attained with the polymer in a molten or melted state, that is above the crystalline melt temperature, or in a dissolved or finely dispersed condition rather than in a solid mass or particulate form. The molten or
25 melted form is more preferred to insure homogeneity rather than localized concentrations at the surface.

While any equipment is suitably used, preferably equipment which provides sufficient mixing and temperature control in the same equipment, but advantageously practice of the invention takes place
30 in such devices as an extruder or a static polymer mixing device such as a Brabender blender. The term extruder is used for its broadest meaning to include such devices as a device which extrudes pellets or pelletizer.

In a preferred embodiment of the invention, the process of the
35 present invention takes place in a single vessel, that is mixing of the coupling agent and polymer takes place in the same vessel as heating to the decomposition temperature of the coupling agent and the foaming process. The vessel is preferably an extruder suitable for foam preparation. The reaction vessel more preferably has at
40 least two zones of different temperatures into which a reaction mixture passes, the first zone advantageously being at a temperature at least the crystalline melt temperature or the softening temperature of the polymer(s) and preferably less than the

decomposition temperature of the coupling agents and the second zone being at a temperature sufficient for decomposition of the coupling agent. The first zone is preferably at a temperature sufficiently high to soften the polymer and allow it to combine with the coupling agent through distributive mixing to a substantially uniform admixture. Addition of a blowing agent conveniently occurs in either of these zones, depending on the temperatures advantageous for its use.

For polymers that have softening points above the coupling agent decomposition temperature (preferably greater than 200°C), and especially when incorporation of a lower melting polymer (such as in a concentrate) is undesirable, the preferred embodiment for incorporation of coupling agent is to solution blend the coupling agent in solution or admixture into the polymer, to allow the polymer to imbibe (absorb or adsorb at least some of the coupling agent), and then to evaporate the solvent. After evaporation, the resulting mixture is extruded. The solvent is preferably a solvent for the coupling agent, and more preferably also for the polymer when the polymer is soluble such as in the case of polycarbonate. Such solvents include polar solvents such as acetone, THF (tetrahydrofuran) and chlorinated hydrocarbons such as methylene chloride. Alternatively other non-polar compounds such as mineral oils in which the coupling agent is sufficiently miscible to disperse the coupling agent in a polymer, are used.

Practice of the process of the invention to rheology modify polymer blends and form foams thereof yields rheology modified or chain coupled polymer foams, that is foams of polymers which have sulfonamide, amine, alkyl-substituted or aryl-substituted carboxamide, alkyl-substituted or aryl-substituted phosphoramidate, alkyl-substituted or aryl-substituted methylene coupling between different polymer chains. Such coupled polymer blends advantageously show higher low shear viscosity than the original polymer due to coupling of long polymer chains to polymer backbones. Broad molecular weight distribution polymers (polydispersity (P.D.) of 3.5 and greater) and gel levels less than 10 percent as determined by xylene extraction show less improvement than the dramatic effect noted in narrow MWD polymer (P.D. = 2.0) with gel less than 10 percent as determined by xylene extraction. The latter are, therefore, preferred for use in the practice of the invention. Coupling leads to polymers which have controlled rheological properties, specifically improved melt strength as evidenced by increased low shear viscosity.

Foam forming steps of the process are within the skill in the art. For instance as exemplified by the excellent teachings to processes for making ethylenic polymer foam structures and processing them in C. P. Park. "Polyolefin Foam", Chapter 9, Handbook of Polymer
5 Foams and Technology, edited by D. Klempner and K. C. Frisch, Hanser Publishers, Munich, Vienna, New York, Barcelona (1991), which is incorporated here in by reference.

The resulting foam structure is optionally made by a conventional extrusion foaming process. The structure is
10 advantageously prepared by heating an ethylenic polymer material to form a plasticized or melt polymer material, incorporating therein a blowing agent to form a foamable gel, and extruding the gel through a die to form the foam product. Prior to mixing with the blowing agent, the polymer material is heated to a temperature at or above its glass
15 transition temperature or melting point. The blowing agent is optionally incorporated or mixed into the melt polymer material by any means known in the art such as with an extruder, mixer, blender, or the like. The blowing agent is mixed with the melt polymer material at an elevated pressure sufficient to prevent substantial
20 expansion of the melt polymer material and to advantageously disperse the blowing agent homogeneously therein. Optionally, a nucleator is optionally blended in the polymer melt or dry blended with the polymer material prior to plasticizing or melting. The foamable gel is typically cooled to a lower temperature to optimize physical
25 characteristics of the foam structure. The gel is then extruded or conveyed through a die of desired shape to a zone of reduced or lower pressure to form the foam structure. The zone of lower pressure is at a pressure lower than that in which the foamable gel is maintained prior to extrusion through the die. The lower pressure is optionally
30 superatmospheric or subatmospheric (vacuum), but is preferably at an atmospheric level.

In another embodiment, the resulting foam structure is optionally formed in a coalesced strand form by extrusion of the ethylenic polymer material through a multi-orifice die. The orifices
35 are arranged so that contact between adjacent streams of the molten extrudate occurs during the foaming process and the contacting surfaces adhere to one another with sufficient adhesion to result in a unitary foam structure. The streams of molten extrudate exiting the die take the form of strands or profiles, which desirably foam,
40 coalesce, and adhere to one another to form a unitary structure. Desirably, the coalesced individual strands or profiles should remain adhered in a unitary structure to prevent strand delamination under stresses encountered in preparing, shaping, and using the foam.

Apparatuses and method for producing foam structures in coalesced strand form are seen in U.S. Pat. Nos. 3,573,152 and 4,824,720.

Alternatively, the resulting foam structure is conveniently formed by an accumulating extrusion process as seen in U.S. Pat. No. 4,323,528. In this process, low density foam structures having large lateral cross-sectional areas are prepared by: 1) forming under pressure a gel of the ethylenic polymer material and a blowing agent at a temperature at which the viscosity of the gel is sufficient to retain the blowing agent when the gel is allowed to expand; 2) extruding the gel into a holding zone maintained at a temperature and pressure which does not allow the gel to foam, the holding zone having an outlet die defining an orifice opening into a zone of lower pressure at which the gel foams, and an openable gate closing the die orifice; 3) periodically opening the gate; 4) substantially concurrently applying mechanical pressure by a movable ram on the gel to eject it from the holding zone through the die orifice into the zone of lower pressure, at a rate greater than that at which substantial foaming in the die orifice occurs and less than that at which substantial irregularities in cross-sectional area or shape occurs; and 5) permitting the ejected gel to expand unrestrained in at least one dimension to produce the foam structure.

In another embodiment, the resulting foam structure is formed into non-crosslinked foam beads suitable for molding into articles. To make the foam beads, discrete resin particles such as granulated resin pellets are: suspended in a liquid medium in which they are substantially insoluble such as water; impregnated with a blowing agent by introducing the blowing agent into the liquid medium at an elevated pressure and temperature in an autoclave or other pressure vessel; and rapidly discharged into the atmosphere or a region of reduced pressure to expand to form the foam beads. This process is well taught in U.S. Pat. Nos. 4,379,859 and 4,464,484.

One modification of the uncrosslinked bead process, styrene monomer is optionally impregnated into the suspended pellets prior to their impregnation with blowing agent to form a graft interpolymer with the ethylenic polymer material. The polyethylene/polystyrene interpolymer beads are cooled and discharged from the vessel substantially unexpanded. The beads are then expanded and molded by an expanded polystyrene bead molding process within the skill in the art. A process of making polyethylene/polystyrene interpolymer beads is described for instance in U.S. Pat. No. 4,168,353.

The foam beads are conveniently then molded by any means within the skill in the art, such as charging the foam beads to the mold, compressing the mold to compress the beads, and heating the beads

such as with steam to effect coalescing and welding of the beads to form the article. Optionally, the beads are impregnated with air or other blowing agent at an elevated pressure and temperature prior to charging to the mold. Further, the beads are optionally heated prior to charging. The foam beads are conveniently then molded to blocks or shaped articles by a suitable molding method within the skill in the art such as taught for instance in U.S. Pat. Nos. 3,504,068 and 3,953,558. Excellent teachings of the above processes and molding methods are seen in C.P. Park, *supra*, p. 191, pp. 197-198, and pp. 227-229.

Blowing agents useful in making the resulting foam structure include inorganic agents, organic blowing agents and chemical blowing agents. Suitable inorganic blowing agents include carbon dioxide, nitrogen, argon, water, air, nitrogen, and helium. Organic blowing agents include aliphatic hydrocarbons having 1-6 carbon atoms, aliphatic alcohols having 1-3 carbon atoms, and fully and partially halogenated aliphatic hydrocarbons having 1-4 carbon atoms. Aliphatic hydrocarbons include methane, ethane, propane, n-butane, isobutane, n-pentane, isopentane, and neopentane. Aliphatic alcohols include methanol, ethanol, n-propanol, and isopropanol. Fully and partially halogenated aliphatic hydrocarbons include fluorocarbons, chlorocarbons, and chlorofluorocarbons. Examples of fluorocarbons include methyl fluoride, perfluoromethane, ethyl fluoride, 1,1-difluoroethane (HFC-152a), 1,1,1-trifluoroethane (HFC-143a), 1,1,1,2-tetrafluoro-ethane (HFC-134a), pentafluoroethane, difluoromethane, perfluoroethane, 2,2-difluoropropane, 1,1,1-trifluoropropane, perfluoropropane, dichloropropane, difluoropropane, perfluorobutane, perfluorocyclobutane. Partially halogenated chlorocarbons and chlorofluorocarbons for use in this invention include methyl chloride, methylene chloride, ethyl chloride, 1,1,1-trichloroethane, 1,1-dichloro-1 fluoroethane (HCFC-141b), 1-chloro 1,1-difluoroethane (HCFC-142b), 1-dichloro-2,2,2-trifluoroethane (HCFC-123) and 1-chloro-1,2,2,2-tetrafluoroethane (HCFC-124). Fully halogenated chlorofluorocarbons include trichloromonofluoromethane (CFC-11), dichlorodifluoromethane (CFC-12), trichlorotrifluoroethane (CFC-113), 1,1,1-trifluoroethane, pentafluoroethane, dichlorotetrafluoroethane (CFC-114), chloroheptafluoropropane, and dichlorohexafluoropropane. Chemical blowing agents include azodicarbonamide, azodiisobutyronitrile, barium azodicarboxylate, N,N'-dimethyl-N,N'-dinitrosoterephthalamide, and benzenesulfonylhydrazide, 4,4-oxybenzene sulfonyl semicarbazide, and p-toluene sulfonyl semicarbazide trihydrazino triazine. Preferred blowing agents include isobutane, HFC-152a, and mixtures of the foregoing.

The amount of blowing agent incorporated into the polymer melt material to make a foam-forming polymer gel is from 0.2 to 5.0, preferably from 0.5 to 3.0, and most preferably from 1.0 to 2.50 gram moles per kilogram of polymer.

5 Foams are optionally perforated to enhance or accelerate permeation of blowing agent from the foam and air into the foam. The foams are optionally perforated to form channels which extend entirely through the foam from one surface to another or partially through the foam. The channels are advantageously spaced up to 2.5
10 centimeters apart and preferably up to 1.3 centimeters apart. The channels are advantageously present over substantially an entire surface of the foam and preferably are uniformly dispersed over the surface. The foams optionally employ a stability control agent of the type described above in combination with perforation to allow
15 accelerated permeation or release of blowing agent while maintaining a dimensionally stable foam. Such perforation is within the skill in the art, for instance as taught in U.S. Patent Nos. 5,424,016 and 5,585,058.

 Various additives are optionally incorporated in the resulting
20 foam structure such as stability control agents, nucleating agents, inorganic fillers, pigments, antioxidants, acid scavengers, ultraviolet absorbers, Flame retardants, processing aids, and extrusion aids.

 A stability control agent is optionally added to the present
25 foam to enhance dimensional stability. Preferred agents include amides and esters of C10-24 fatty acids. Such agents are seen in U.S. Pat. Nos. 3,644,230 and 4,214,054. Most preferred agents include stearyl stearamide, glyceromonostearate, glycerol monobehenate, and sorbitol monostearate. Typically, such stability control agents are
30 employed in an amount ranging from 0.1 to 10 parts per hundred parts of the polymer.

 The resulting foam structure preferably exhibits excellent dimensional stability. Preferred foams recover 80 or more percent of initial volume within a month with initial volume being measured
35 within 30 seconds after foam expansion. Volume is measured by a suitable method such as cubic displacement of water.

 In addition, a nucleating agent is optionally added in order to control the size of foam cells. Preferred nucleating agents include inorganic substances such as calcium carbonate, talc, clay, titanium
40 oxide, silica, barium sulfate, diatomaceous earth, mixtures of citric acid and sodium bicarbonate. The amount of nucleating agent employed may range from 0.01 to 5 parts by weight per hundred parts by weight of a polymer resin.

The resulting foam structure is substantially noncrosslinked or uncrosslinked. The polymer material comprising the foam structure is substantially free of crosslinking. The foam structure contains no more than 5 percent gel as measured according to ASTM D-2765-84

- 5 Method A. A slight degree of crosslinking, which occurs naturally without the use of crosslinking agents or radiation, is permissible.

The resulting foam structure preferably has a density of less than 250, more preferably less than 100 and most preferably from 10 to 70 kilograms per cubic meter. The foam preferably has an average
10 cell size of from 0.05 to 5.0, more preferably from 0.2 to 2.0, and most preferably 0.3 to 1.8 millimeters as measured according to the procedures of ASTM D3576.

The resulting foam structure optionally is in any physical configuration within the skill in the art, such as extruded sheet,
15 rod, plank, and profiles. The foam structure is optionally formed by molding for example of expandable beads into any of the foregoing configurations or any other configuration.

The resulting foam structure is optionally closed-celled or open-celled. Preferably, the present foam contains 80 percent or more
20 closed cells as measured according to ASTM D2856-A.

In the most preferred embodiment the foam forming step or steps and the coupling steps are at least partially simultaneous. Thus the coupling agent is introduced during any step before or in a foam forming process that is of a temperature sufficiently low to result
25 in adequate mixing before or during coupling and coupling takes place in or simultaneous with any step in a foam forming process in which the temperature is at least the decomposition temperature of the coupling agent. Coupling, however, preferably takes place before the foam is extruded or otherwise exits the vessel in which the polymer
30 is admixed with any blowing agent.

Foams prepared according to the practice of the invention advantageously have one or preferably more of smaller cell diameter, homogeneous cell diameter distribution, lower foam density, higher tensile or compressive strength, or higher tensile or compressive
35 toughness (or combinations thereof) than foams made from the same starting materials but not coupled using C-H insertion coupling agents, preferably sulfonyl azides, more preferably than foams formed from the same starting materials and coupled using free radical means, especially peroxides. Cell size is measured according to the
40 procedure of ASTM D-3576; tear strength is measured according to the procedure of ASTM D-624; tensile properties including tensile strength, compressive strength and toughness are measured according to the procedures of ASTM D-412. Additives such as antioxidants (for

example, hindered phenols such as, for example, Irganox™ 1010), phosphites (for example, Irgafos™ 168) both commercially available from Ciba Geigy Corporation), U. V. stabilizers, cling additives (for example, polyisobutylene), antiblock additives, colorants, pigments, and fillers, are optionally also included in the interpolymers employed in the blends of the present invention, to the extent that they do not interfere with the enhanced properties discovered by Applicants.

The additives are advantageously employed in functionally equivalent amounts known to those skilled in the art. For example, the amount of antioxidant employed is that amount which prevents the polymer or polymer blend from undergoing oxidation at the temperatures and environment employed during storage and ultimate use of the polymers. Such amount of antioxidants is usually in the range of from 0.01 to 10, preferably from 0.05 to 5, more preferably from 0.1 to 2 percent by weight based upon the weight of the polymer or polymer blend. Similarly, the amounts of any of the other enumerated additives are the functionally equivalent amounts such as the amount to render the polymer or polymer blend antiblocking, to produce the desired amount of filler loading to produce the desired result, to provide the desired color from the colorant or pigment. Such additives are advantageously employed in the range of from 0.05 to 50, preferably from 0.1 to 35, more preferably from 0.2 to 20 percent by weight based upon the weight of the polymer or polymer blend. Fillers, however are advantageously employed in amounts up to 90 percent by weight based on the weight of the polymer or polymer blend.

The blends of the present invention, in addition to the production of foams, are advantageously used to produce a wide range of fabricated articles including calendared, cast and blown sheets and films, compression and injection molded parts, fibers. The blends are also useful in applications such as modifiers for bitumen and asphalt compositions and as components for hot melt and pressure sensitive adhesive systems.

The following examples are to illustrate this invention and do not limit it. Ratios, parts, and percentages are by weight unless otherwise stated. Examples (Ex) of the invention are designated numerically while comparative samples (C.S.) are designated alphabetically and are not examples of the invention.

Test Methods

The viscosity of polymer as a function of shear rate was measured according to the following method. A dynamic mechanical spectrometer commercially available from Rheometrics, Inc. under the trade designation RMS-800 with 25 mm diameter parallel plates was used to
5 determine the dynamic rheological data. A frequency sweep with five logarithmically spaced points per decade was run from 0.1 to 100 rad/s at 190 °C. The strain was determined to be within the linear viscoelastic regime by performing a strain sweep at 0.1 rad/s and 190 °C, by strain sweep from 2 to 30 percent strain in 2 percent steps to
10 determine the minimum required strain to produce torques within the specification of the transducer; another strain sweep at 100 rad/s and 190°C was used to determine the maximum strain before nonlinearity occurred according to the procedure disclosed by J. M. Dealy and K. F. Wissbrun, "Melt Rheology and Its Role in Plastics
15 Processing", Van Nostrand, New York (1990).

Viscosity and melt index tests were performed in a nitrogen purge to minimize oxidative degradation.

20 A thermomechanical analyzer (TMA) commercially available from Perkin Elmer Corporation under the trade designation model TMA 7 was used to measure the upper service temperature (UST). Probe force of 102g and heating rate of 5°C/min were used. Each test specimen was a disk with thickness of 2mm and diameter, prepared by compression molding
25 at 205°C and air-cooling to room temperature. Temperature at the probe penetration of 1 mm is taken as the upper service temperature.

Xylene Extraction to determine gel content was performed by weighing out 1 gram samples of polymer. The samples are transferred to a mesh
30 basket which is then placed in boiling xylene for 12 hours. After 12 hours, the sample baskets are removed and placed in a vacuum oven at 150°C and 28 in. of Hg vacuum for 12 hours. After 12 hours, the samples are removed, allowed to cool to room temperature over a 1 hour period, and then weighed. The results are reported as percent
35 polymer extracted. Percent extracted = (initial weight-final weight)/initial weight according to ASTM D-2765 Procedure "A"

Tensile properties were determined by compression molding 1/16 inch plaques. Tensile specimens were then cut from these plaques and
40 tested on an instrument commercially available from Instron Corporation under the trade designation Instron model 1122 load

frame using 0.870 inch (2.2 cm) micro-tensile samples measured at an extension rate of 5 inch/min (12.7 cm/min). Tensile at break and elongation at break, were measured in accordance with ASTM D-412. The toughness was measured as the area under the stress/strain curve.

5

The melt index for the ethylene based polymers was measured according to ASTM D-1238 under conditions of 190°C/2.16 Kg (formerly known as Condition E).

- 10 General procedures for determining compression set are described in ASTM D 395-89. The sample plaques were cut into disks of 1.14 inch (2.90 cm) diameter. The disks were stacked up to a thickness of 0.5 inch (1.27 cm). Test specimens were measured under constant strain of 25 percent, at 70°C for 22 h. The sample was aged at 70°C for 22
15 h under 25 percent compression, then cooled to 22°C.

Test parts and characterization data for the interpolymers and their blends are generated according to the following procedures:

- Compression Molding: Samples are melted at 190°C for 3 minutes
20 and compression molded at 190°C under 20,000 lb (9,072 kg) of pressure for another 2 minutes. Subsequently, the molten materials are quenched in a press equilibrated at room temperature.

Density: The density of the samples is measured according to ASTM-D792.

25

The following materials were used:

- PE (polyethylene):** An ethylene octene copolymer with melt index of 1 g/10 min, density of 0.902 g/cc commercially available from Dow
30 Chemical Company under the trade designation Affinity PL1880 polyolefin plastomer is used as the polyethylene.

- PP (polypropylene):** Isotactic polypropylene used in the blends is commercially available from Himont Incorporated under the trade designation Profax 6523 polypropylene. The product has a melt flow
35 rate of 4.0 g/10 min (measured at 230°C). This product has a Tensile strength at yield of 5000 PSI (34500kPa) and a flex modulus of 250,000 PSI (1723,500 kPa).

- HDPE:** HDPE is a high density polyethylene with melt index of 30 g/min, density of 0.9600 g/10 min., commercially available from Dow
40 Chemical Company under the trade designation HDPE HD 30460 M polyethylene.

ESI-1: an experimental ethylene/styrene copolymer made using a solution process for which procedural detail is included hereafter.. This copolymer is 41 weight percent styrene and 58 weight percent ethylene. The overall sample also contains 2.0 weight percent
5 homopolymer styrene as an impurity. It has a melt index (MI) of 0.1 g/10 minutes at 190°C under 2.16 kg load.

ESI -2: is an ethylene -styrene interpolymers having 60 weight percent of styrene and 40 weight percent of ethylene, containing atactic polystyrene of 7.5 weight percent, and having a melt index of 0.5
10 g/10 min.

The Ethylene Styrene Interpolymers (ESI-1 and ESI-2) are synthesized according to the following general procedure:

Reactor Description

15 A 6 gallon (22.7 L), oil jacketed, autoclave continuously stirred tank reactor (CSTR) was employed as the reactor. A magnetically coupled agitator with impellers commercially available from Lightning Mixers, Inc. under the trade designation A-320
20 impellers provides the mixing. The reactor ran liquid full at 475 psig (3,275 kPa). Process flow was in the bottom and out the top. A heat transfer oil was circulated through the jacket of the reactor to remove some of the heat of reaction. After the exit from the reactor there was a flow meter that measured flow and solution density. All
25 lines on the exit of the reactor were traced with 50 psi (344.7 kPa) steam and insulated.

Procedure

Solvent (ethylbenzene for ESI-1 and toluene for ESI-2) was supplied to the reactor at 30 psig (207 kPa). The feed to the
30 reactor was measured by a mass flow meter. A variable speed diaphragm pump controlled the feed rate of solvent. At the discharge of the solvent pump, a side stream was taken to provide flush flows for the catalyst injection line (1 lb/hr (0.45 kg/hr)) and the reactor agitator (0.75 lb/hr (0.34 kg/hr)). These flows were
35 measured by differential pressure flow meters and controlled by manual adjustment of micro-flow needle valves. Uninhibited styrene monomer was supplied to the reactor at 30 psig (308 kPa). The feed to the reactor was measured by a mass flow meter. A variable speed diaphragm pump controlled the feed rate. The styrene stream was
40 mixed with the remaining solvent stream. Ethylene was supplied to the reactor at 600 psig (4,238 kPa). The ethylene stream was measured by a mass flow meter just prior to a valve controlling flow. A flow meter controller was used to deliver hydrogen into the

ethylene stream at the outlet of the ethylene control valve. The ethylene/hydrogen mixture combines with the solvent/styrene stream at ambient temperature. The temperature of the solvent/monomer as it enters the reactor was reduced to 5°C by a heat exchanger with -5°C glycol on the jacket thereof. This solvent/styrene stream entered the bottom of the reactor. The three component catalyst system described in Table 2 and its solvent flush also enter the reactor at the bottom but through a different port than the monomer stream. Preparation of the catalyst components took place in an inert atmosphere glove box. The diluted components were put in nitrogen padded cylinders and charged to catalyst run tanks for the reaction. From these run tanks the catalyst was pressured with piston pumps and the flow was measured with flow meters. These streams combine with each other and the catalyst flush solvent just prior to entry through a single injection line into the reactor where they react to form the designated polymer.

Polymerization was stopped as reaction mixture flows into a reactor product line after the reactor, by addition of catalyst kill (water mixed with solvent) into the reactor product line after a flow meter which measures solution density. A static mixer in the line provided dispersion of the catalyst kill and additives in the reactor effluent stream. This stream next entered post reactor heaters that provide additional energy for the solvent removal flash. This flash occurred as the effluent exited the post reactor heater and the pressure was dropped from 475 psig (3,275 kPa) down to ~250mm Hg (33 kPa) of pressure absolute at the reactor pressure control valve. This flashed polymer entered a hot oil jacketed devolatilizer. Approximately 85 percent of the volatile compounds (hereinafter volatiles) were removed from the polymer in the devolatilizer. The volatiles exit the top of the devolatilizer. The stream of exiting volatiles was condensed and with a glycol jacketed exchanger, entered the suction of a vacuum pump and was discharged to a glycol jacket solvent and styrene/ethylene separation vessel. Solvent and styrene were removed from the bottom of the vessel and ethylene from the top. The ethylene stream was measured with a flow meter and analyzed for composition. The measurement of vented ethylene plus a calculation of the dissolved gasses in the solvent/styrene stream were used to calculate the ethylene conversion. The polymer separated in the devolatilizer was pumped out with a gear pump to an extruder commercially available from Werner Pfleiderer Corporation under the trade designation ZSK-30 devolatilizing vacuum extruder. The dry polymer exits the extruder as a single strand. This strand was cooled as it was pulled through a water bath. The excess water was

blown from the strand with air and the strand was chopped into pellets with a strand chopper.

The catalyst used in preparing ESI-1 was (t-butylamido) dimethyl(tetramethylcyclopentadienyl)silane-titanium (II) 1,3-pentadiene. The catalyst used in preparing ESI 2 was Titanium, [1,1'-(η^4 -1,3-butadiene-1,4-diyl)bis(benzene)][1-[(1,2,3,3a,11b- η)-1H-cyclopenta[1]phenanthren-1-yl]-N-(1,1-dimethylethyl)-1,1-dimethylsilanaminato(2-)-N]-. The cocatalyst was bis-hydrogenated tallowalkyl methylammonium tetrakis(pentafluorophenyl)borate. A modified methylaluminoxane commercially available from Akzo Nobel Chemicals Inc. under the trade designation MMAO-3A was also used in the amounts indicated in Tables 1 and 2 and is referred to herein as MMAO.

Table 2a Catalyst to Cocatalyst ratio and MMAO to Catalyst Ratio

	Molar ratio of cocatalyst/Catalyst	Molar ratio of MMAO to catalyst
ESI-1	1.25:1	10.0:1
ESI-2	5:1	15:1

Table 2b Reactor Data

Polymer	Reactor Temp. °C	Solv. Flow		Ethylene Flow		Hydrogen Flow SCCM	Styrene Flow		Ethylene Conv percent percent
		lb/hr	kg/hr	lb/hr	kg/hr		lb/hr	kg/hr	
ESI-1	94	37	16.76	2.8	0.41	3.0	5.0	2.27	96.5
ESI-2	67.9	30	13.59	1.3	0.186	0	10	4.53	86.3

SCCM means standard cubic centimeter

Conv. means conversion

Solv. means solvent.

Temp. means temperature.

Preparation of 4,4'-disulfonylazidophenyl ether:

4,4'-bis(chlorosulfonyl)phenyl ether (10g, 0.027 mole) was dissolved in 100 mL of acetone and 4.426 g (0.06808 moles) of solid sodium azide was added portionwise over the course of 15 minutes. The reaction mixture was stirred for 26 hours at

ambient temperature and then was filtered to remove sodium chloride. The resulting filter cake was washed with acetone and the combined filtrate evaporated to yield a white solid which was washed twice with 20 mL portions of water and then dried at
5 ambient temperature under vacuum. The resulting white solid (7.3 g, 70 percent yield) was identified as 4,4'-disulfonylazidophenyl ether by ¹H and ¹³C NMR spectroscopy.

Comparative Sample A.

10 In a mixing bowl (40.0g) heated to 190°C with the roller blades turning at 75 RPM was placed 14.0 grams of the polypropylene. After the polypropylene had melted, 26.0 g of ES 41 was added. After addition of the ES 41 polymer, the admixed sample was mixed at 190°C for 10 minutes. The sample was removed and
15 allowed to cool. The sample was compression molded into a 1/16 inch (1.58 mm) thick plaque at 190°C and 20,000 pounds (5,072 kg) of force for 8 minutes. The sample was removed and cooled. The tensile properties, the UST, the compression set at 70°C and the percent gel by xylene extraction were determined. The
20 results are shown in Table 1.

Examples 1 and Comparative Example B

The procedure of Comparative Example A was repeated except that 2 minutes after the addition of the second polymer, 0.08g
25 (0.2mmol, 0.2 weight percent) of 4,4'-disulfonylazidophenyl ether was added. The sample was mixed for 5 more minutes and removed from the mixer. This sample is Example 1.

For Comparative Sample B, procedure of Example 1 was followed
30 except that 0.20g (0.5 mmole, 0.5 weight percent) of 4,4'-disulfonylazidophenyl ether was added.

After cooling, both samples were compression molded into a 1/16 inch (1.58 mm) thick plaque at 190°C and 20,000 pounds
35 (5,072 kg) of force for 8 minutes. The samples were removed and cooled. The tensile properties, the UST, the compression set at 70°C and the percent gel by xylene extraction of the samples were determined. The results are shown in Table 1.

Table 3:

Sample	UST (TMA)	Percent Change in UST	MI (12, 190°C)	MI (110, 190°C)	percent Elongation	Break Stress (PSI)	Break Stress in kPa	Compression Set	percent Gel
C.S. A	98	0	0.4	8.3	520	2900	19990	100percent	0
EX. 1	142	44.9	0.2	15.8	510	2700	18610	100percent	0
C.S. B	160	63.2	<0.01	1.5	350	2300	15870	80percent	53

The examples show how a small amount of (0.2 weight percent) of 4,4'-disulfonylazidophenyl ether is capable of improving the temperature resistance of ethylene styrene interpolymer/polypropylene blends without inducing crosslinking (as evidenced by the 0 percent gel measurement of Example 1) or deterioration of the physical properties. Furthermore, the sample was still melt processable and demonstrated increased processability at higher shear rates (as evidenced by the I10 values). Control Sample B showed that the processability of the blend decreased (low melt index) and gels were formed if excessive sulfonylazide (more than 0.4 weight percent) is used.

The amount of azide used was advantageously adjusted so that the gel was less than 2 percent. High molecular weight of the ethylene styrene interpolymer used requires less azide than is required by Polymers of lower molecular weight.

Visual examination of Transmission Electron Micrographs of samples of the compositions prepared in Example 1 and Comparative Sample A shows that the size of regions of polypropylene phase in the blend with sulfonyl azide was smaller and more continuous than the control sample without sulfonylazide.

Example 2:

A mixer commercially available from Haake Fusion Co. consisting of a mixer with the trade designation HaakeBuchler Rheomix 3000 mixer with roller style blades, attached to a rheometer with the trade designation HaakeBuchler Rheocord 9000 Torque rheometer. The mixing bowl was heated to 126°C. The mixing speed was set at 20 rpm. ESI 2 (152 g) and an ethylene octene copolymer was commercially available from Dow Chemical Company under the trade designation of Affinity PL 1880 polyolefin plastomers. After 2 minutes, 0.30 g of 4,4'-disulfonylazidophenyl (0.15 weight percent) was added to the mixer and mixed for 1 minute. The temperature of the mixture was raised to 190-200°C. This was done by raising the set temperature to 160°C and increasing the stirring rate to 75 rpm (revolutions per minute). After 10 minutes, the sample was removed from the mixer bowl.

C.S. C:

The method of Ex 2 was repeated except that no 4,4'-disulfonylazidophenyl was used.

Example 3:

The method of Ex. 2 was repeated except that 148.6g of ESI-2, 49.5 g of a the HDPE HD30460 M polyethylene, and 0.3g of 4,4'-disulfonylazidophenyl were used.

5

C.S. D:

The method of Ex 2 was repeated except that 148.6g of ESI-2 and 49.5 g of the HDPE HD30460 M polyethylene were used.

10

Table 4. Properties of Azide Modified Ethylene Octene Copolymer and Ethylene Styrene Interpolymer Blend

Sample and Comparative Sample	UST (°C)	0.1 Rad/s Viscosity (poise)	0.1 Rad/s Viscosity (Pa-s)	Ratio of 0.1 to 100 Rad/s Viscosity
C.S. C	84	1.60E05	1.60E04	7.96
Ex. 2	90	13.32E05	13.32E04	65.07
C.S. D	96	N/M	N/M	N/M
EX. 3	110	N/M	N/M	N/M

N/M: not measured

- 15 The results in Table 4 shows the products from blends of alpha-olefin/vinyl aromatic or hindered aliphatic vinyl monomer interpolymer with polyolefins reacted with and not reacted with poly(sulfonyl azide). This result was obtained with blends of ethylene/styrene interpolymers and either an ethylene/octene polymer
- 20 or HDPE. The data from C.S. C and Ex.2 indicate the blend using poly(sulfonyl azide) (Ex.2) had higher melt strength (0.1 rad viscosity) and higher shear thinning effect (ratio of 0.1 rad/s to 10 rad/s viscosity) than the sample without poly(sulfonyl azide) (C.S. C).

What is claimed is:

1. A process of preparing a coupled polymer blend characterized in heating an admixture containing (1) a polymer blend containing:
 - 5 (A) from 1 to 99 weight percent of one or more α -olefin/hindered vinyl monomer substantially random interpolymers, each having been made from monomer components comprising:
 - (1) from 0.5 to 65 mole percent of either
 - (a) at least one vinyl aromatic monomer, or
 - 10 (b) at least one hindered aliphatic vinyl monomer, or
 - (c) a combination of at least one vinyl aromatic monomer and at least one hindered aliphatic vinyl monomer; and
 - (2) from 35 to 99.5 mole percent of at least one aliphatic α -olefin having from 2 to 20 carbon atoms; and
 - 15 (B) from 99 to 1 weight percent based on entire composition of one or more homopolymers or copolymers made from monomer components comprising aliphatic α -olefins having from 2 to 20 carbon atoms, or aliphatic α -olefins having from 2 to 20 carbon atoms and containing polar groups; and (2) a coupling amount of at least one poly(sulfonyl
 - 20 azide) to at least the decomposition temperature of the poly(sulfonyl azide) for a period sufficient for decomposition of at least 80 weight percent of the poly(sulfonyl azide) and sufficient to result in a coupled polymer blend having less than 2 weight percent gel.
2. The process of Claim 1 wherein in the blend: component (A) is a
 - 25 substantially random interpolymer made from styrene and ethylene, or styrene, ethylene and at least one other α -olefin containing from 3 to 8 carbon atoms; component (B) is a homopolymer made from ethylene or propylene, or a copolymer made from ethylene, or propylene or a combination thereof and at least other α -olefin containing from 4 to
 - 30 8 carbon atoms; or a terpolymer of ethylene, propylene and at least one of 4-methyl pentene, butene-1, hexene-1 or octene-1; and the amount of poly(sulfonyl azide) is from 0.01 to 1 weight percent of the blend.
3. The process of Claim 1 wherein the coupling agent comprises at least
 - 35 one poly(sulfonyl azide) which has a structure X-R-X wherein each X is SO_2N_2 , and R represents an unsubstituted or inertly substituted hydrocarbyl, hydrocarbyl ether or silicon-containing group; wherein at least one poly(sulfonyl azide) has at least 3 but less than 50 carbon, silicon or oxygen atoms between sulfonyl azide groups.

4. The process of Claim 4 wherein R includes at least one aryl group between the sulfonyl groups; and wherein R includes at least two aryl groups or wherein R is one aryl group, and the group has more than one ring and wherein the poly(sulfonyl azide) and blend are mixed at
5 a first temperature which is at least the melting point of the lowest melting component of the blend and after mixing react at a second temperature which is at least greater than the first temperature, is at least the decomposition temperature of the poly(sulfonyl azide) and is greater than 185°C.
- 10 5. A composition comprising a reaction product obtainable by any of the processes of Claims 1-4.
6. An article which comprises a composition of Claim 5.
7. The article of Claim 6 wherein the article is formed from a melt of the composition of Claim 5.
- 15 8. The article of Claim 6 or 7 which is a calendared, cast and blown sheet, film, compression and injection molded part, fiber, modifier for bitumen or asphalt compositions, or a component in a hot melt or pressure sensitive adhesive system.
9. A foam formed from a foam forming composition comprising the
20 composition of Claim 5.
10. The process of formation of the article of Claim 6 or 7 by calendering, blowing, casting, injection molding, compression molding, extrusion, blow molding, foaming, or spinning. a composition of Claim 6.

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INTERNATIONAL SEARCH REPORT

International Application No

PCT/US 98/17668

A. CLASSIFICATION OF SUBJECT MATTER

IPC 6 C08K5/43 C08L23/16 C08L23/02

According to International Patent Classification (IPC) or to both national classification and IPC

B. FIELDS SEARCHED

Minimum documentation searched (classification system followed by classification symbols)

IPC 6 C08K C08G C08F C08J C08L

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched

Electronic data base consulted during the international search (name of data base and, where practical, search terms used)

C. DOCUMENTS CONSIDERED TO BE RELEVANT

Category *	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
Y	DE 15 69 025 A (HERCULES) 9 July 1970 cited in the application see examples 25,26 ---	1-10
A	US 3 203 936 A (D.S.BRESLOW ET AL) 31 August 1965 see column 1, line 26 - column 2, line 25 ---	1-10
A	US 3 855 184 A (BOSTICK E ET AL) 17 December 1974 see example 1 ---	1-10
Y	WO 96 07681 A (DOW CHEMICAL CO) 14 March 1996 see page 12, line 33 - line 36 -----	1-10



Further documents are listed in the continuation of box C.



Patent family members are listed in annex.

* Special categories of cited documents :

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Date of the actual completion of the international search

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INTERNATIONAL SEARCH REPORT

Information on patent family members

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